



Supplement of

Soil bacterial community and functional shifts in response to altered snowpack in moist acidic tundra of northern Alaska

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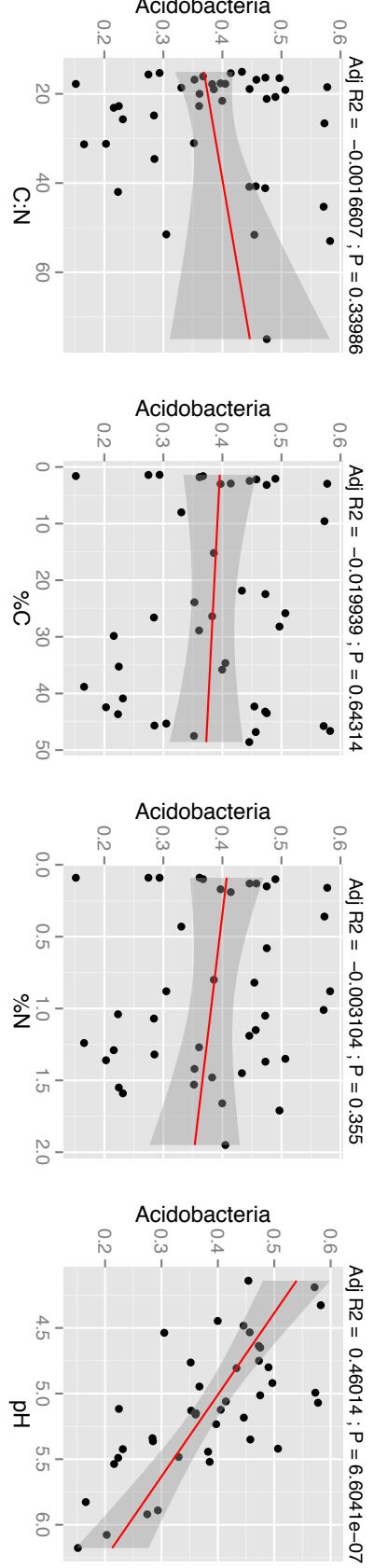
3 **Table S1** - List of targeted KEGG ortholog enzymes used in the PICRUST analyses. For more information see materials and methods
4 in main text.

Supplemental Information

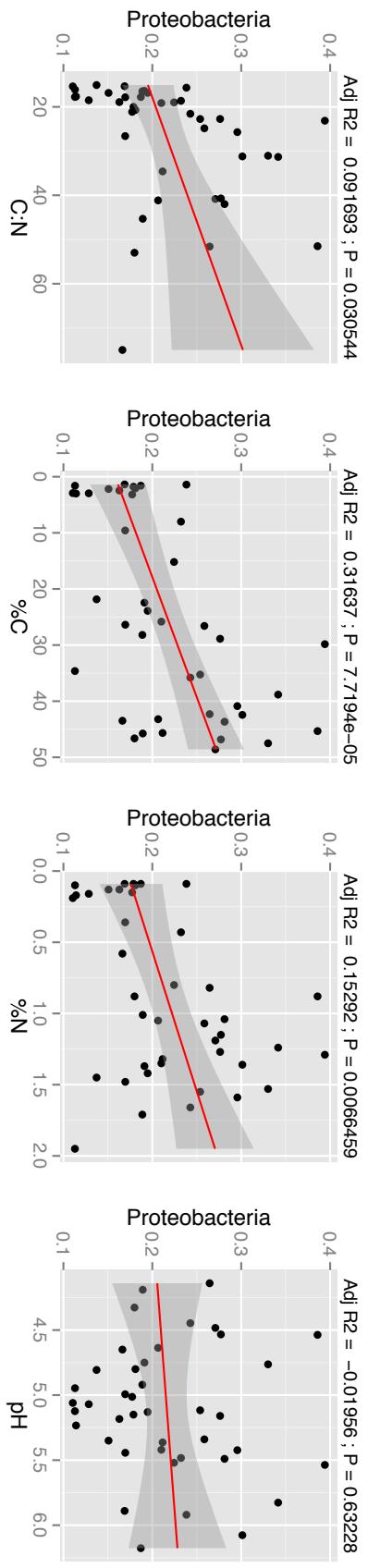
Functional role	Enzyme name	EC number	KEGG orthology number
PRESENT IN SAMPLES			
Arabinoside degradation	arabinogalactan endo-beta-1,4-galactanase arabinan endo-1,5-alpha-L-arabinanase	3.2.1.89 3.2.1.99	K01224 K06113
Cellulose degradation	cellulase beta-glucosidase cellulose 1,4-beta-cellulobiosidase (CBH)	3.2.1.4 3.2.1.21 3.2.1.91	K01179 K01188, K05349, K05350 K01225
Chitin degradation	chitinase (NAG) bifunctional chitinase/lysozyme	3.2.1.14 3.2.1.14 & 3.2.1.17	K01183 K13381
Nitrogen mobilization	leucyl aminopeptidase (LAP) urease	3.4.11.1 3.5.1.5	K01255 K01427, K01428, K01429, K01430, K14048
Lignin degradation	tyrosinase (phenol oxidase)	1.14.18.1	K00505
Pectin degradation	polygalacturonase alpha-L-thamnosidase	3.2.1.15 3.2.1.40	K01184 K05989
Phosphate mobilization	alkaline phosphatase (AP) acid phosphatase (AP)	3.1.3.1 3.1.3.2	K01077, K01113 K01078, K01093, K09474, K03788, K14379
Superoxides	superoxide dismutase superoxide reductase	1.15.1.1 1.15.1.2	K00518, K04564, K04565 K05919
Xylan degradation	xylan 1,4-beta-xylosidase	3.2.1.37	K01198
NOT FOUND IN ANY SAMPLES			
	polyphenol oxidase	1.10.3.1	K00422

peroxidase	1.11.1.7	K00430
fructan beta-fructosidase	3.2.1.80	K03332
laccase (oxidoreductase)	1.10.3.2	K05909
eosinophil peroxidase	1.11.1.7	K10788
cytosol aminopeptidase	3.4.11.1 & 3.4.11.5	K11142
peroxiredoxin 6, 1-Cys peroxiredoxin	1.11.1.7, 1.11.1.15, & 3.1.1,-	K11188
lactoperoxidase	1.11.1.7	K12550
low molecular weight phosphotyrosine protein phosphatase	3.1.3.2 3.1.3.48	K14394
lysophosphatidic acid phosphatase type 6	3.1.3.2	K14395
acid phosphatase	3.1.3.2	K14410
beta-D-xylosidase 4	3.2.1.37	K15920

1 **Figures S1-to-S6.** Linear fit regressions of bacterial phyla abundance with respect to soil properties such as soil carbon-to-nitrogen
 2 ratio (C/N), soil carbon mass concentration (%C), soil nitrogen mass concentration (%N), and soil pH across all snow treatment sites
 3 and all soil depths. The bacterial relative abundance is ordered from most abundant phylum (Fig. S1) to least abundant (Fig S6). See
 4 table S2 for statistics and significance.



5 Fig. S1 – Linear relationships between Acidobacteria relative abundance and soil C:N, C and N concentrations, and pH from all 41
 6 samples. Shaded areas indicate 95% confidence intervals.
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8 Fig. S2 – Linear relationships between Proteobacteria relative abundance and soil C:N, C and N concentrations, and pH from all 41
 9 samples. Shaded areas indicate 95% confidence intervals.
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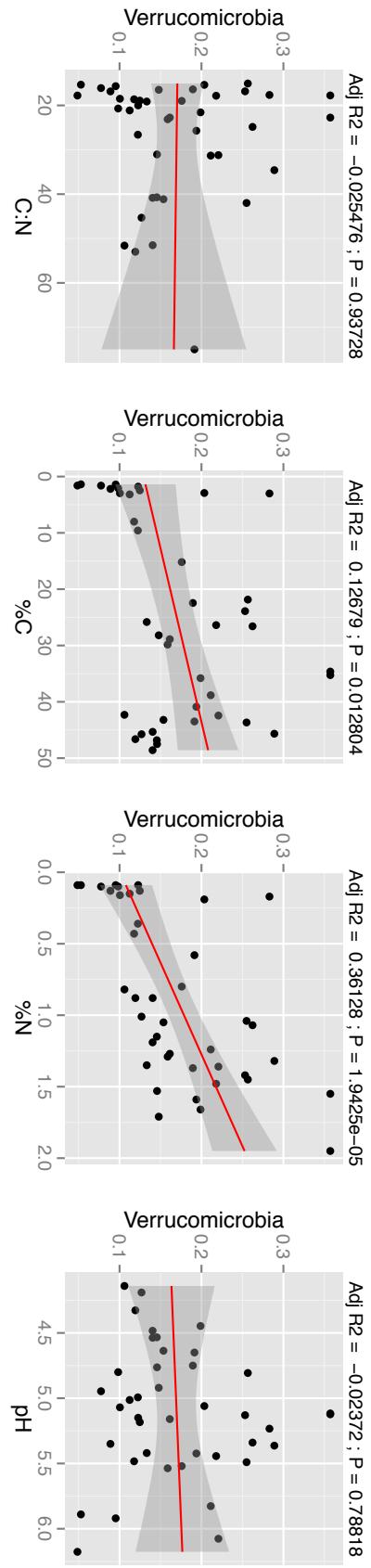


Fig. S3 – Linear relationships between Verrucomicrobia relative abundance and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.

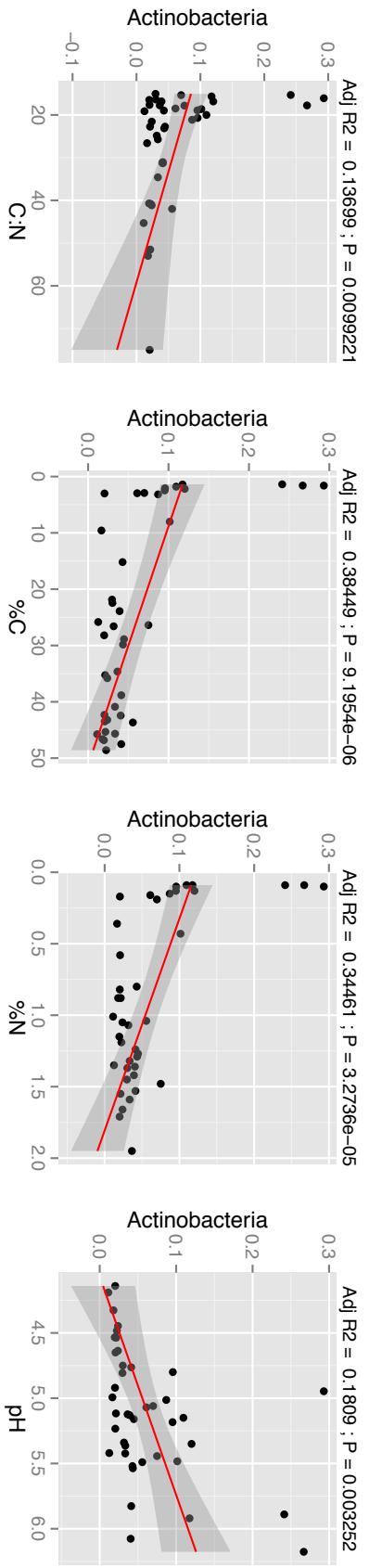


Fig. S4 – Linear relationships between Actinobacteria relative abundance and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.

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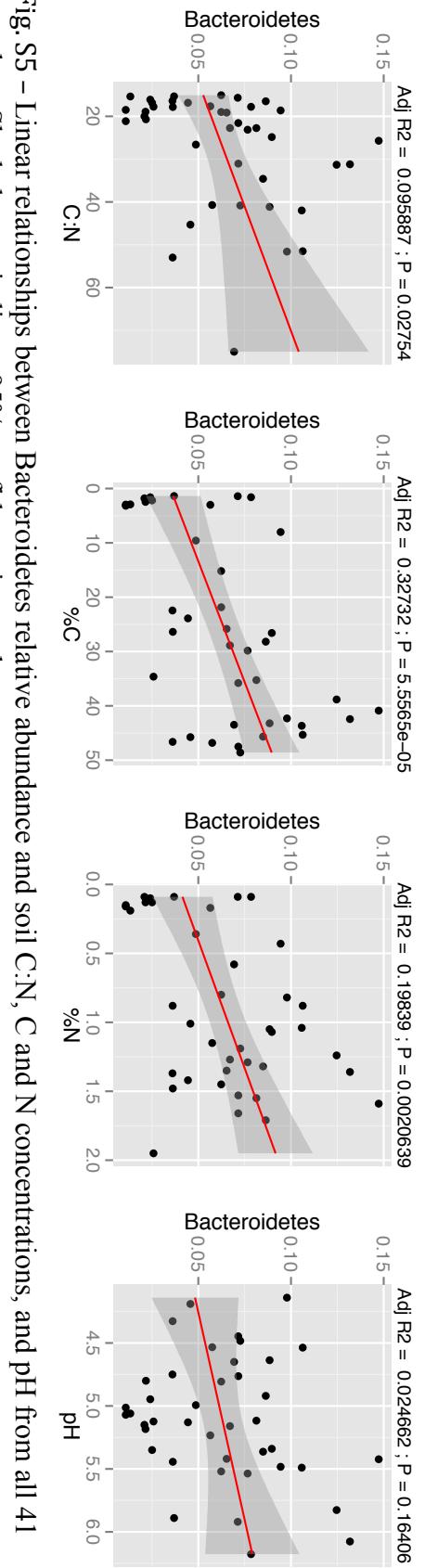


Fig. S5 – Linear relationships between Bacteroidetes relative abundance and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.

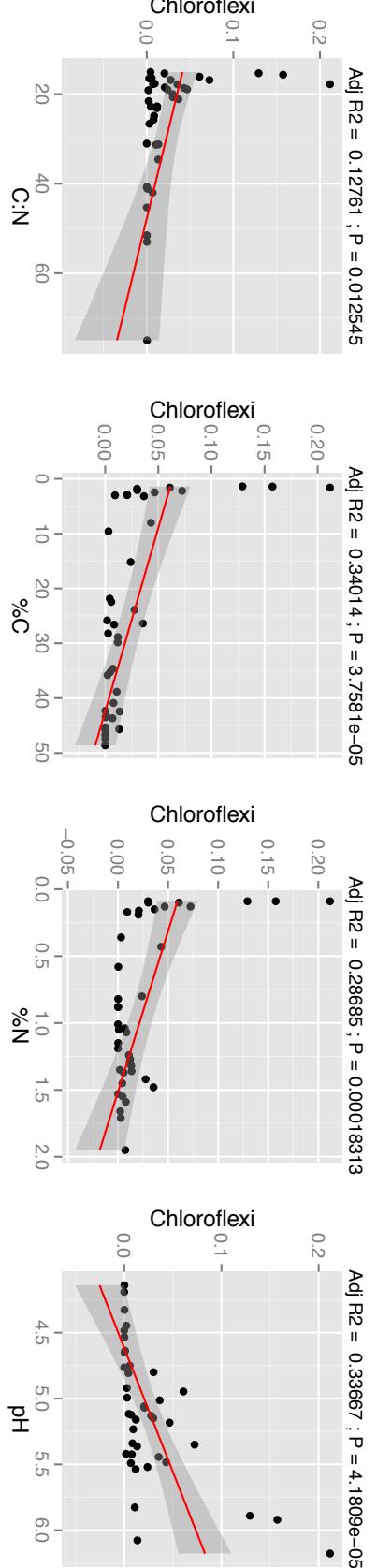
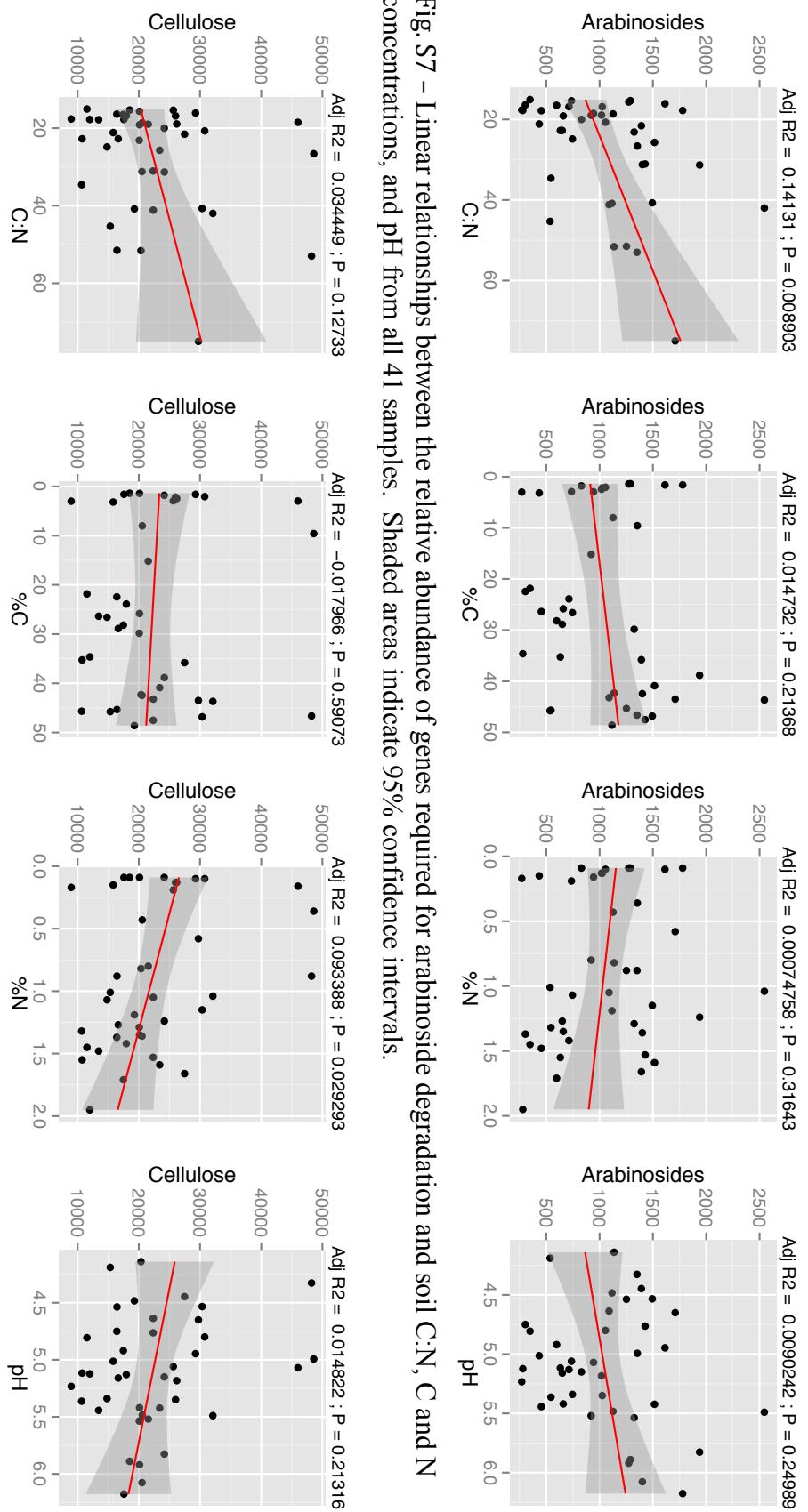


Fig. S6 – Linear relationships between Chloroflexi relative abundance and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.

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Figures S7-to-S15. Linear fit regressions of relative gene copy abundance with respect to soil properties such as soil carbon-to-nitrogen ratio (C/N), soil carbon mass concentration (%C), soil nitrogen mass concentration (%N), and soil pH across all snow treatment sites and all soil depths. Enzymes were grouped by their general activities as explained in main text and they are shown on the Y-axis of the figures. See table S2 for statistics and significance.



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6 Fig. S7 – Linear relationships between the relative abundance of genes required for arabinoside degradation and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.
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9 Fig. S8 – Linear relationships between the relative abundance of genes required for cellulose degradation and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.
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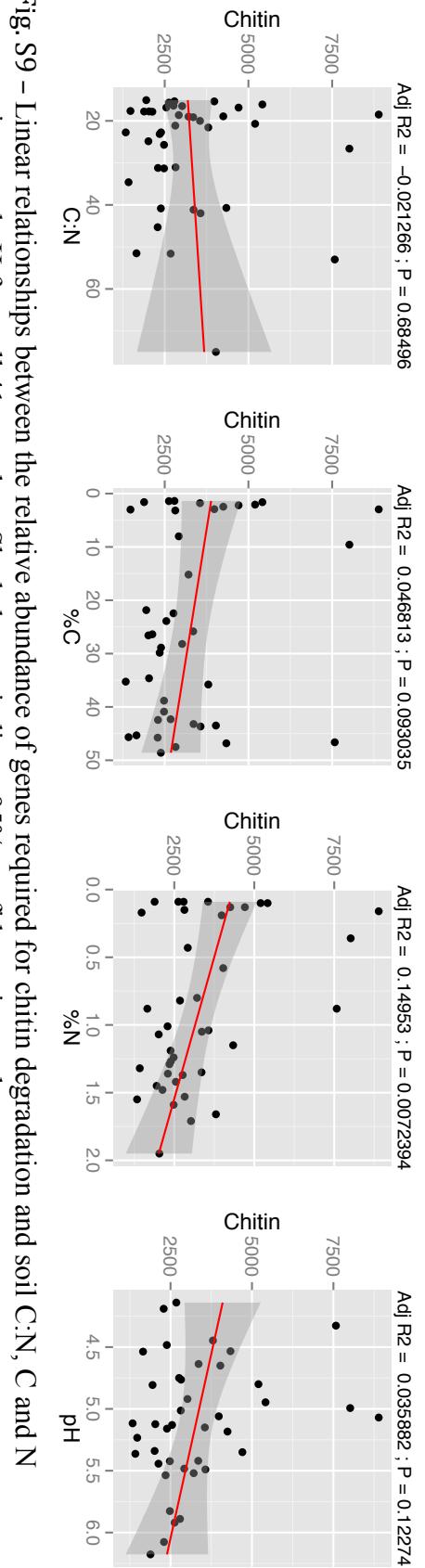


Fig. S9 – Linear relationships between the relative abundance of genes required for chitin degradation and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.

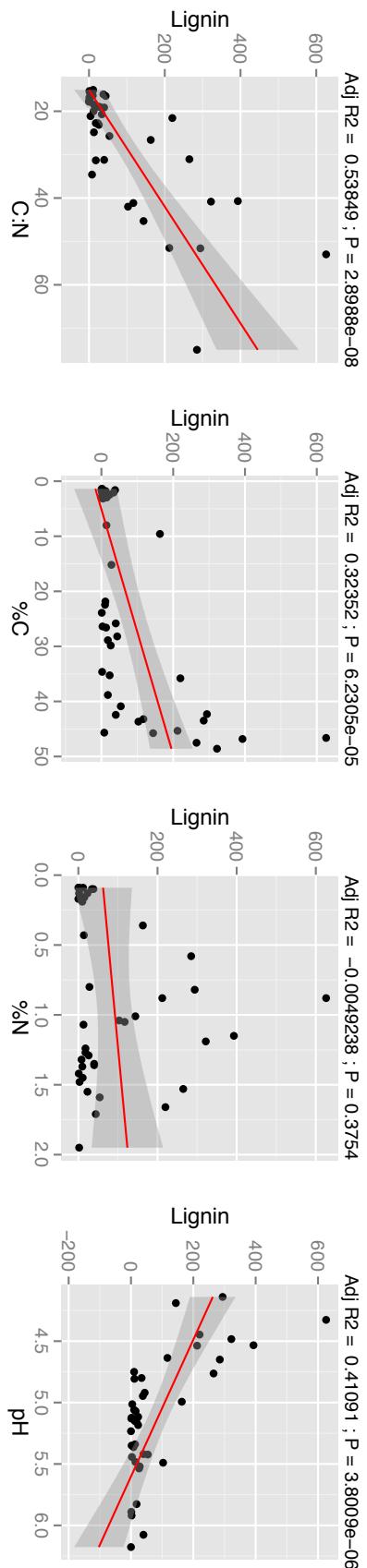
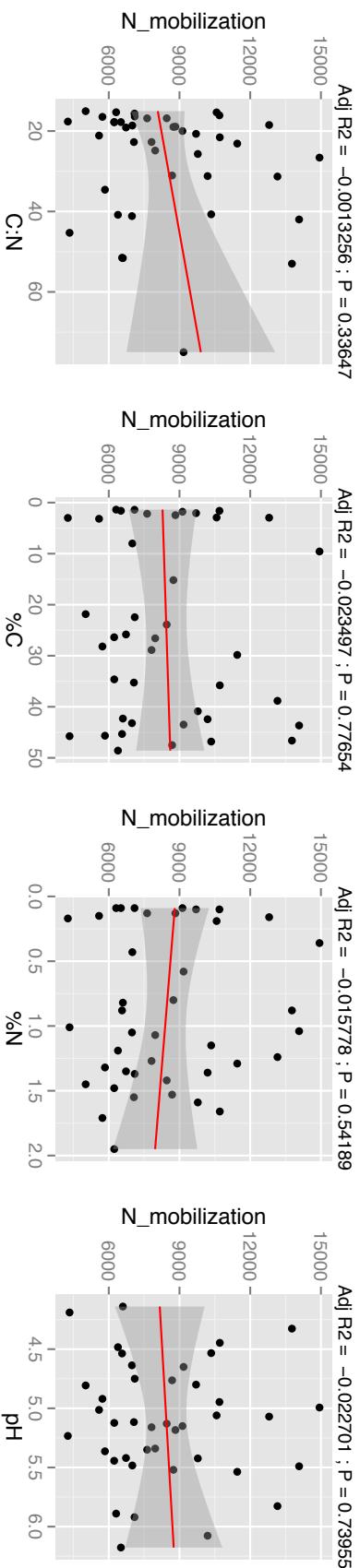
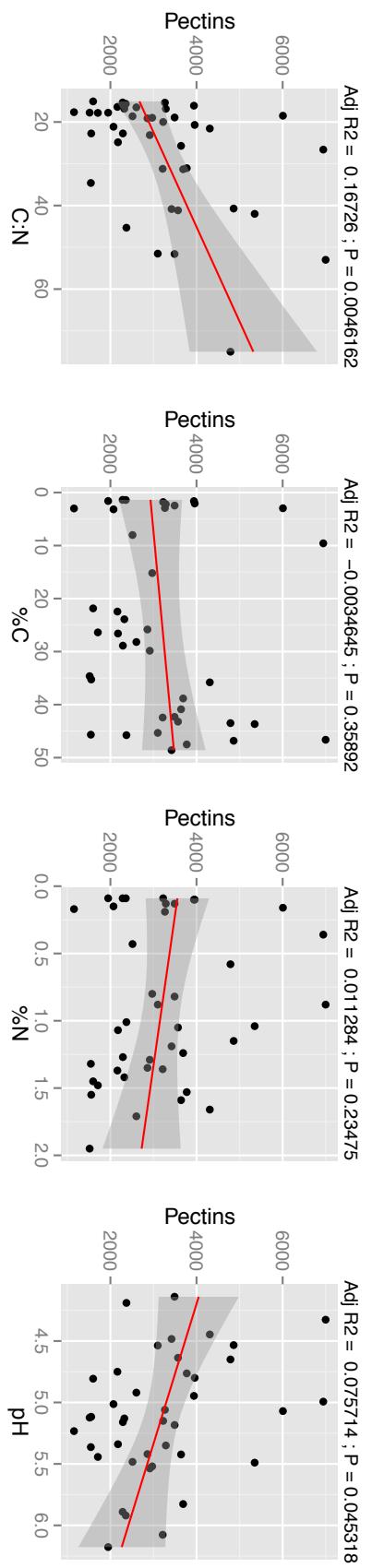


Fig. S10 – Linear relationships between the relative abundance of genes required for lignin degradation and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.

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1 Fig. S11 – Linear relationships between the relative abundance of genes required for N mobilization and soil C:N, C and N
2 concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.
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6 Fig. S12 – Linear relationships between the relative abundance of genes required for pectin degradation and soil C:N, C and N
7 concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.
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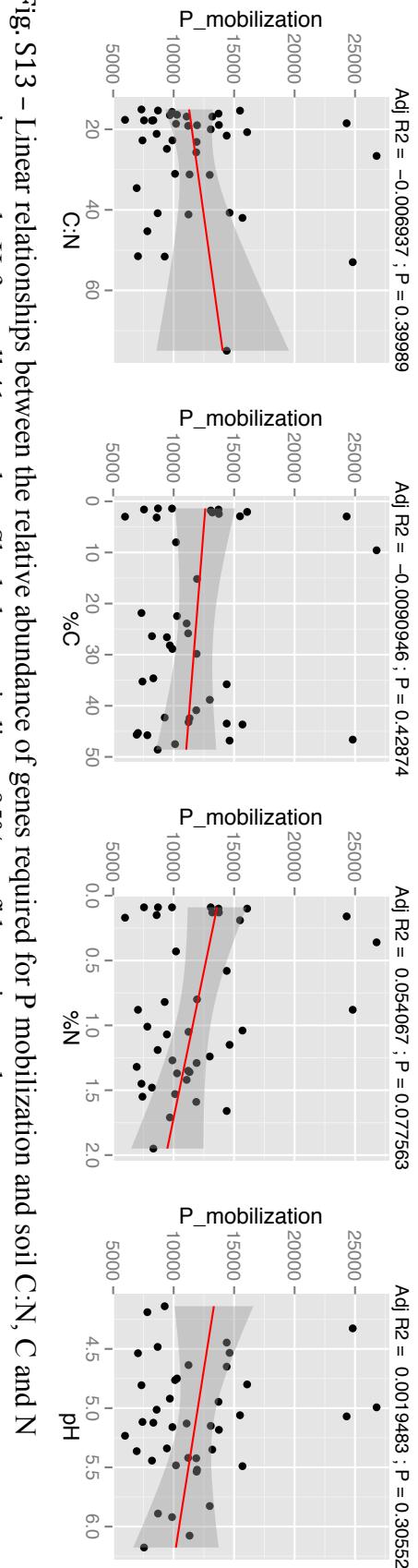


Fig. S13 – Linear relationships between the relative abundance of genes required for P mobilization and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.

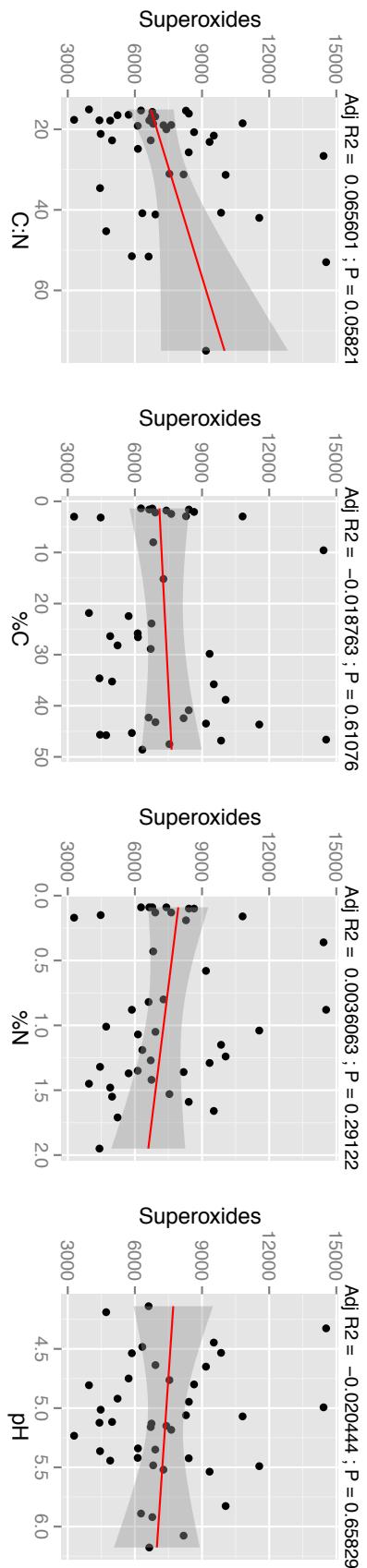


Fig. S14 – Linear relationships between the relative abundance of genes required for superoxide regulation and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.

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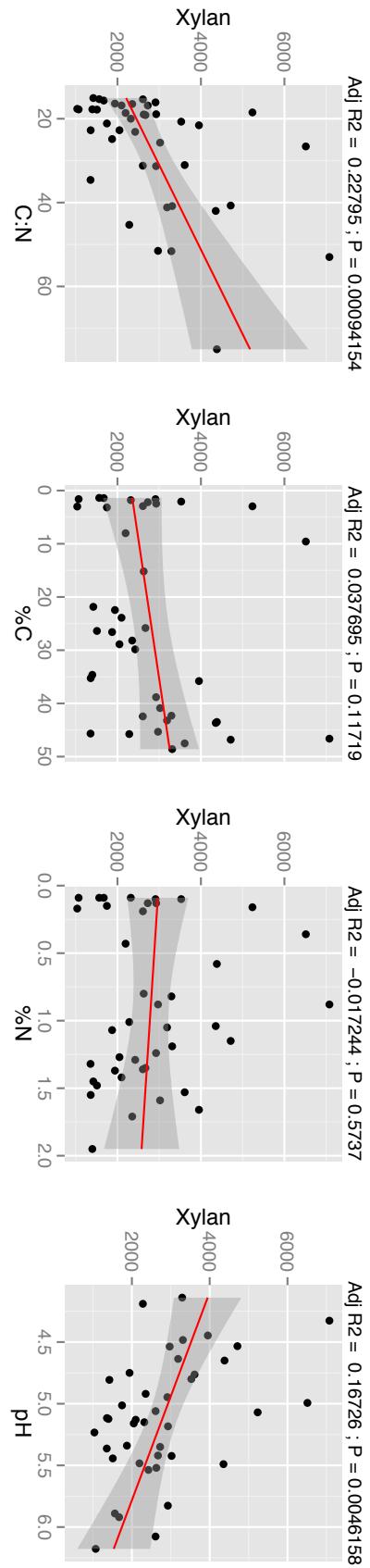


Fig. S15 – Linear relationships between the relative abundance of genes required for xylan degradation and soil C:N, C and N concentrations, and pH from all 41 samples. Shaded areas indicate 95% confidence intervals.

1 Table S2 – Effects of soil depth characteristics (Organic, Transition, Mineral) on soil
 2 chemistry, bacterial phylum relative abundance, and relative abundance of genes organized
 3 by functional groups, as determined by the Kruskal-Wallis test. Degrees of freedom = 2 for
 4 all analyses. * = p<0.05, ** = p<0.01, *** = p<0.001

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	H-statistic	p-value	Soil layer w/ higher value (Org vs Min)
<u>Soil Chemistry</u>			
%C	32.32	9.57 x 10 ⁻⁸ ***	Organic
%N	26.53	1.74 x 10 ⁻⁶ ***	Organic
C:N	21.70	1.94 x 10 ⁻⁵ ***	Organic
pH	6.81	0.03 *	Mineral
<u>Bacterial phylum abundance</u>			
Acidobacteria	0.05	0.98	NA
Proteobacteria	14.78	6.17 x 10 ⁻⁴ ***	Organic
Verrucomicrobia	14.93	5.73 x 10 ⁻⁴ ***	Organic
Actinobacteria	20.16	4.12 x 10 ⁻⁵ ***	Mineral
Bacteroidetes	13.08	1.45 x 10 ⁻³ **	Organic
Chloroflexi	24.80	4.13 x 10 ⁻⁶ ***	Mineral
<u>Enzyme gene abundance</u>			
Lignin	20.17	4.17 x 10 ⁻⁵ ***	Organic
Chitin	3.00	0.22	NA
Cellulose	8.36	0.02 *	Organic
Pectin	17.25	1.80 x 10 ⁻⁴ ***	Organic
Xylan	15.79	3.72 x 10 ⁻⁴ ***	Organic
Arabinoside	13.62	1.10 x 10 ⁻³ **	Organic
N mobilization	11.67	2.92 x 10 ⁻³ **	Organic
P mobilization	3.30	0.19	NA
Superoxide	29.61	3.72 x 10 ⁻⁷ ***	Organic

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1 Table S3 – Statistical analyses of alpha diversity using non-parametric two-sample t-tests
2 with 999 Monte Carlo permutations. Pairwise comparisons of Shannon diversity metrics
3 from each sample were made between each soil layer (Organic, Transition, Mineral) and
4 each snow accumulation treatment (Control, Deep, Intermediate, Low).

5 * = p<0.05, ** = p<0.01, *** = p<0.001.

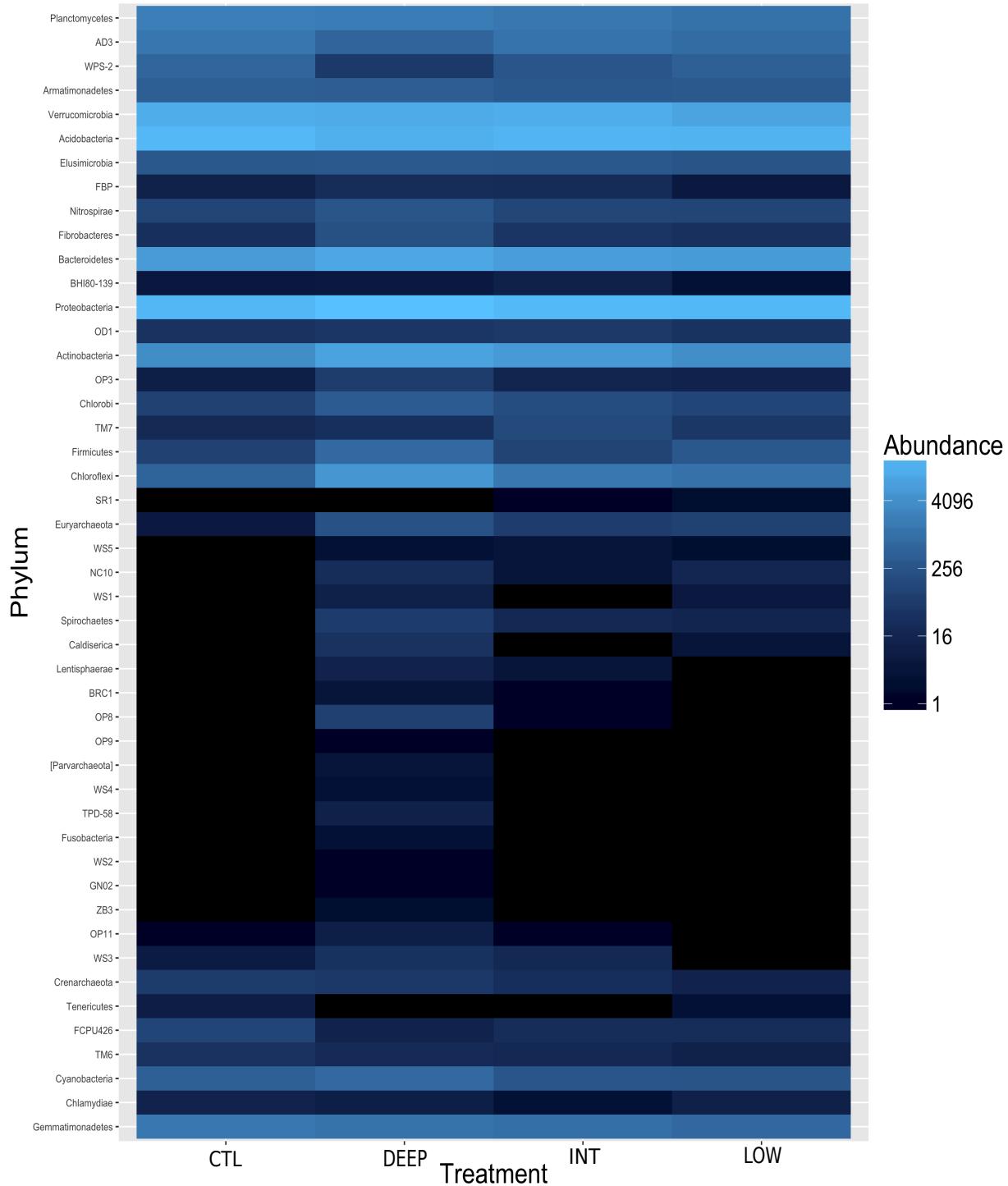
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Two-sample t-tests		
<u>Soil layers</u>	<u>t- statistic</u>	<u>p-value</u>
Organic / Mineral	5.58	0.003 **
Trans / Organic	-0.26	1
Trans / Mineral	5.22	0.003**
<u>Treatment</u>		
Cont / Deep	0.30	1
Cont/ Int	-0.49	1
Cont / Low	1.70	0.6
Deep / Int	-4.0×10^{-4}	1
Deep / Low	-.30	1
Int / Low	0.51	1

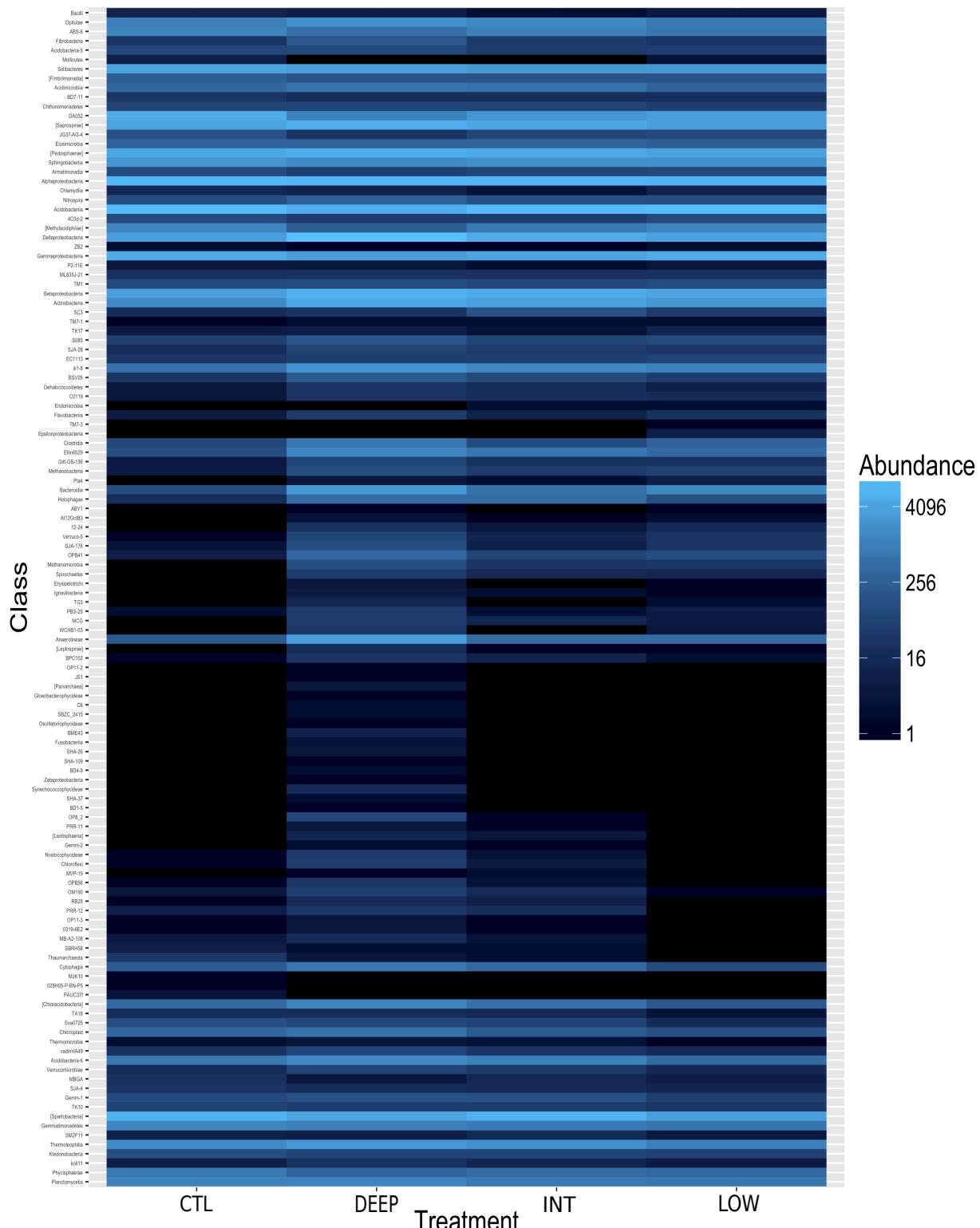
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1 **Figures S16-S20.** Heatmaps showing the relative raw abundance of operational taxonomic
 2 units (OTU) at different taxonomic levels (phyla, class, family, individual OTU) for each of
 3 the snow treatments: control (CTL), 100% more snow accumulation (DEEP), 50% more
 4 snow (INT), and 25% less snow than control (LOW).



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 6 Fig. S16 – Heatmap of raw gene abundance (# of OTU's) for all detected Phyla. Columns
 7 represent snow accumulation treatment groups.

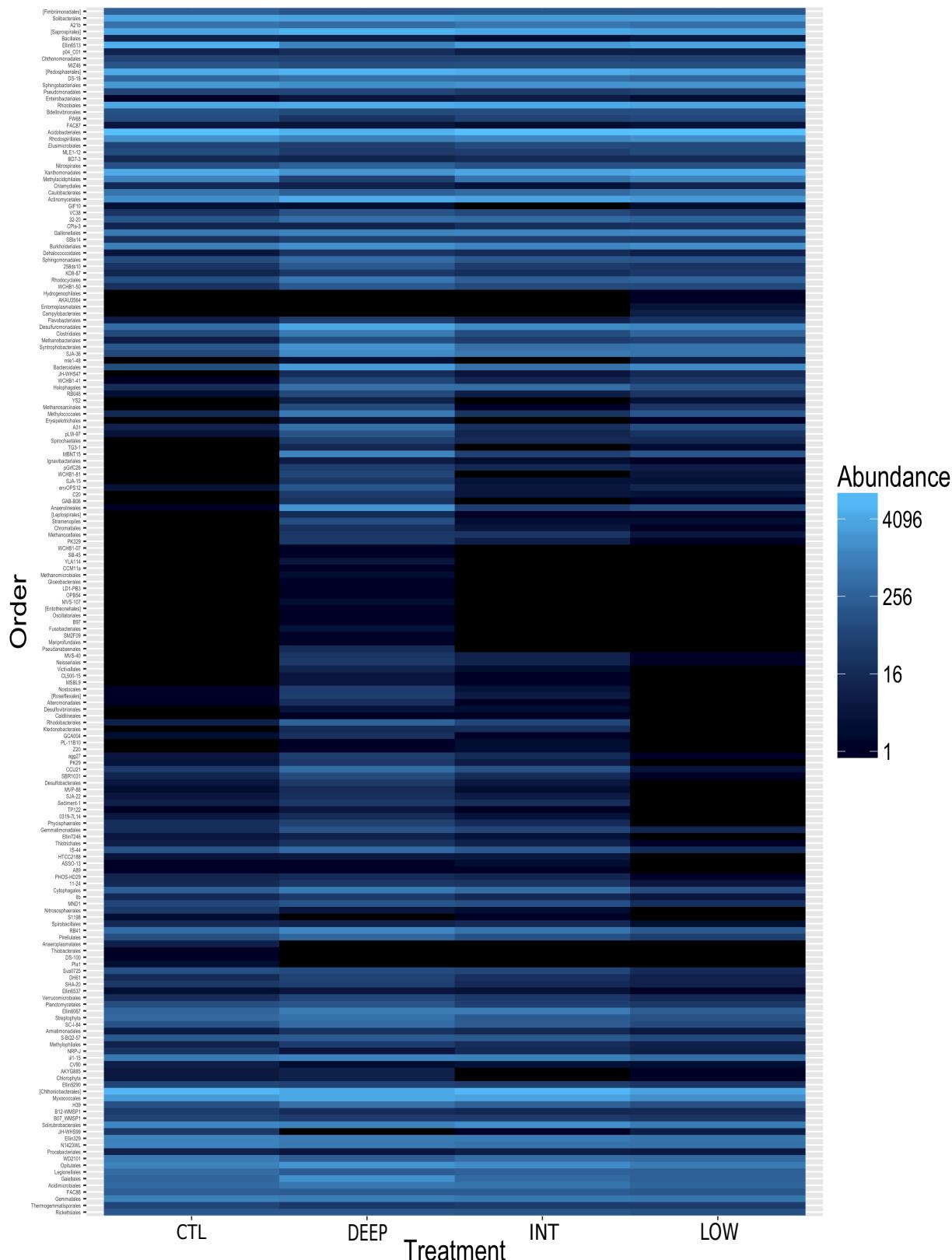


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2 Fig. S17 - Heatmap of raw gene abundance (# of OTU's) for all detected Classes. Columns

3 represent snow accumulation treatment groups.

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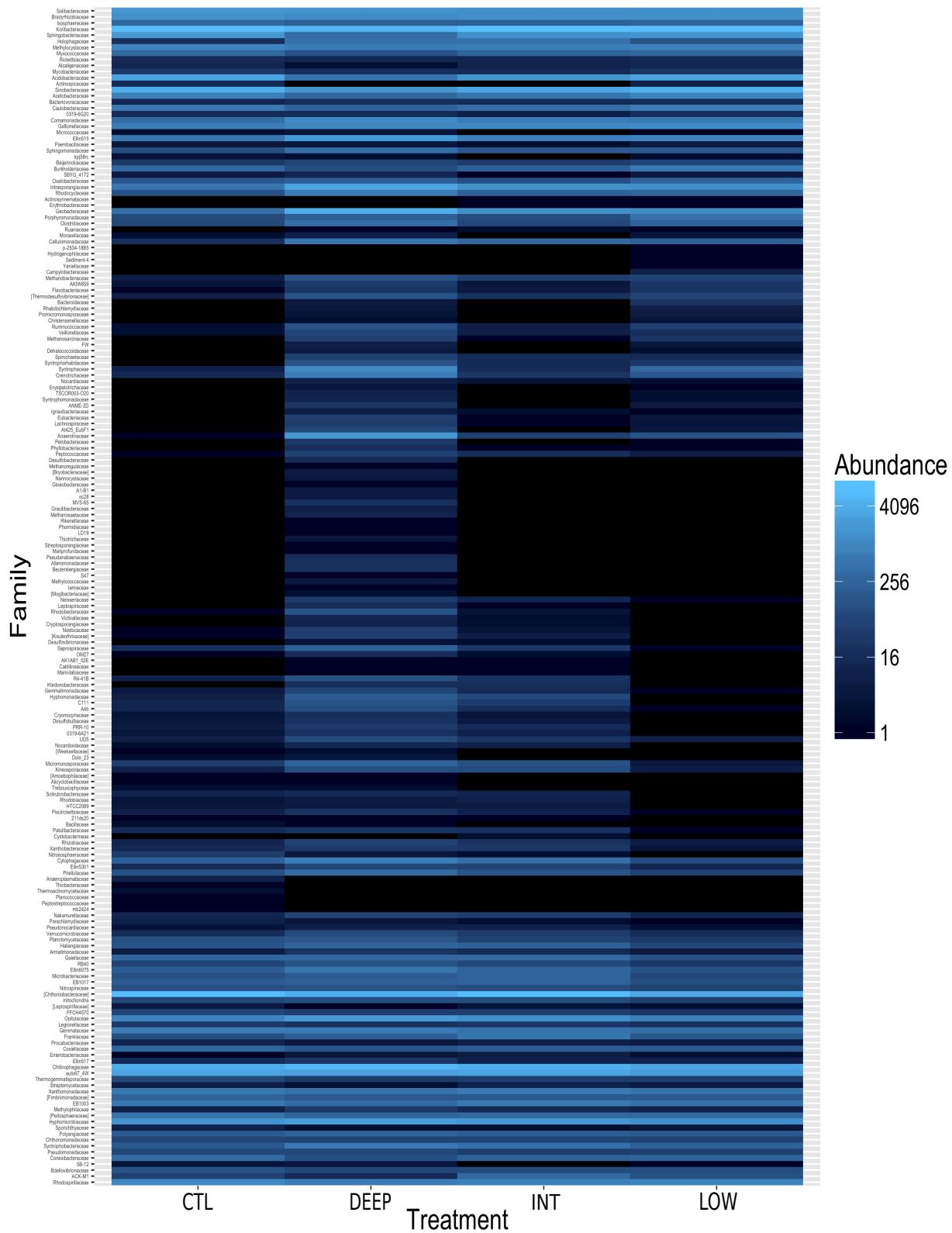
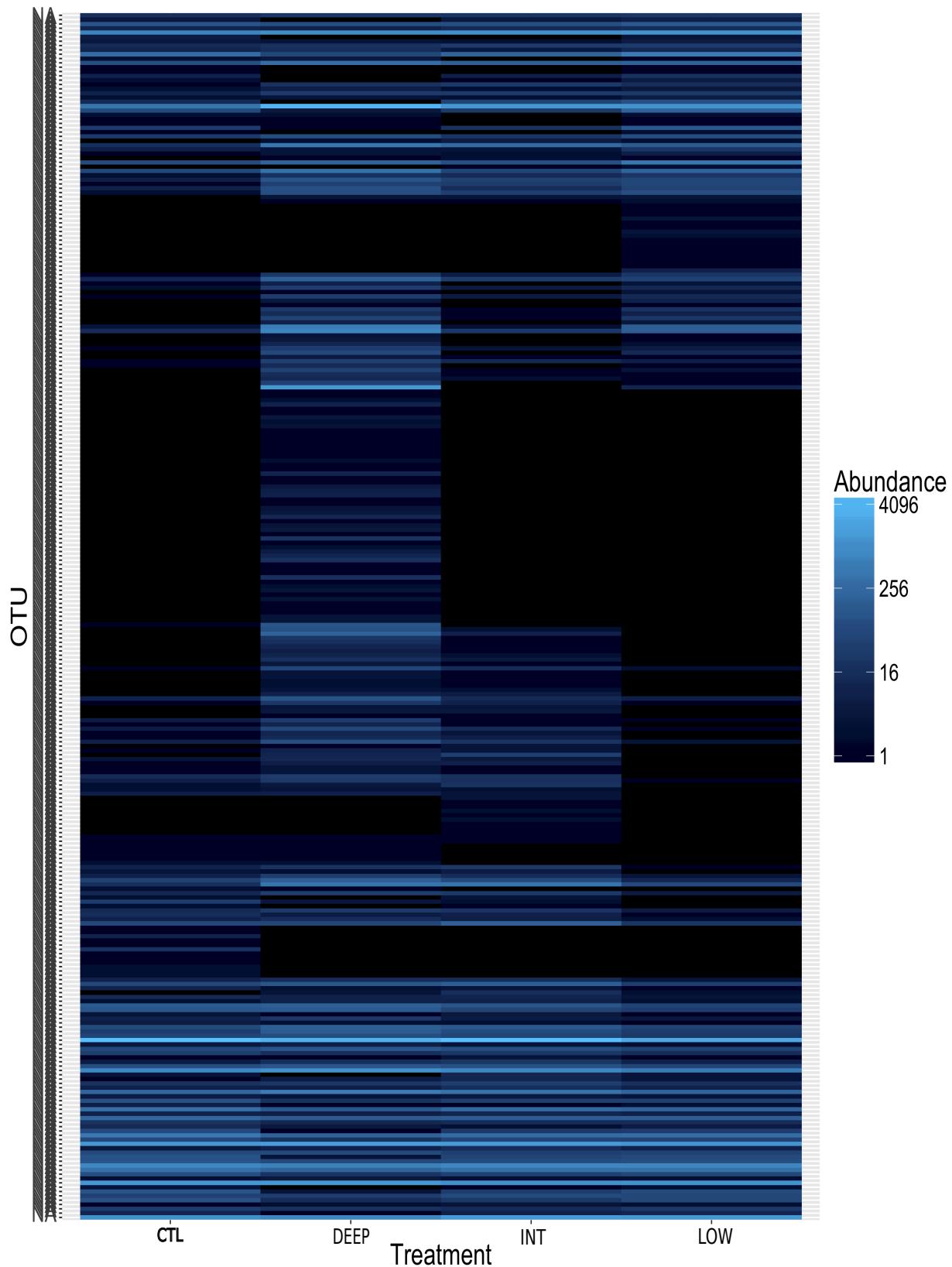


Fig. S19 - Heatmap of raw gene abundance (# of OTU's) for all detected Families. Columns represent snow accumulation treatment groups.



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2 Fig. S20 - Heatmap of raw gene abundance (# of OTU's) for individual OTU assignments.
3 Columns represent snow accumulation treatment groups.